

AD _____

Award Number: DAMD17-01-1-0154

TITLE: Integrin-Mediated Stimulation of HIF-1x and Angiogenesis
in Breast Cancer

PRINCIPAL INVESTIGATOR: Arthur M. Mercurio, Ph.D.

CONTRACTING ORGANIZATION: Beth Israel Deaconess Medical Center
Boston, Massachusetts 02215

REPORT DATE: June 2003

TYPE OF REPORT: Annual

PREPARED FOR: U.S. Army Medical Research and Materiel Command
Fort Detrick, Maryland 21702-5012

DISTRIBUTION STATEMENT: Approved for Public Release;
Distribution Unlimited

The views, opinions and/or findings contained in this report are those of the author(s) and should not be construed as an official Department of the Army position, policy or decision unless so designated by other documentation.

20031106 043

REPORT DOCUMENTATION PAGEForm Approved
OMB No. 074-0188

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing this collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302, and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503

1. AGENCY USE ONLY
(Leave blank)**2. REPORT DATE**
June 2003**3. REPORT TYPE AND DATES COVERED**
Annual (1 Jun 2002 - 31 May 2003)**4. TITLE AND SUBTITLE**Integrin-Mediated Stimulation of HIF-1 α and Angiogenesis in Breast Cancer**5. FUNDING NUMBERS**

DAMD17-01-1-0154

6. AUTHOR(S)

Arthur M. Mercurio, Ph.D.

7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES)Beth Israel Deaconess Medical Center
Boston, Massachusetts 02215

E-Mail: amercuri@caregroup.harvard.edu

**8. PERFORMING ORGANIZATION
REPORT NUMBER****9. SPONSORING / MONITORING****AGENCY NAME(S) AND ADDRESS(ES)**U.S. Army Medical Research and Materiel Command
Fort Detrick, Maryland 21702-5012**10. SPONSORING / MONITORING
AGENCY REPORT NUMBER****11. SUPPLEMENTARY NOTES****12a. DISTRIBUTION / AVAILABILITY STATEMENT**

Approved for Public Release; Distribution Unlimited

12b. DISTRIBUTION CODE**13. ABSTRACT (Maximum 200 Words)**

Understanding the mechanisms involved in the progression from carcinoma *in situ* to metastatic disease is one of the most important problems in breast cancer research. To this end, studies by our group and others have implicated a critical role for the $\alpha 6$ integrins in breast cancer progression. For this reason, it is essential to elucidate the mechanisms by which these integrins contribute to breast cancer progression. This IDEA Award is based on the hypothesis that the $\alpha 6$ integrins regulate expression of VEGF, a growth factor that is essential for tumor survival and angiogenesis. In Year 2 of this Award, we have made considerable progress in validating this hypothesis and we have established that the $\alpha 6 \beta 1$ integrin is necessary for the activation of HIF-1 α , a transcription factor that regulates VEGF expression in breast carcinoma cells.

14. SUBJECT TERMSBreast, angiogenesis, integrins, HIF-1 α **15. NUMBER OF PAGES**

7

16. PRICE CODE**17. SECURITY CLASSIFICATION
OF REPORT**

Unclassified

**18. SECURITY CLASSIFICATION
OF THIS PAGE**

Unclassified

**19. SECURITY CLASSIFICATION
OF ABSTRACT**

Unclassified

20. LIMITATION OF ABSTRACT

Unlimited

NSN 7540-01-280-5500

Standard Form 298 (Rev. 2-89)
Prescribed by ANSI Std. Z39-18
298-102

Table of Contents

Cover.....	1
SF 298.....	2
Table of Contents.....	3
Introduction.....	4
Body.....	4
Key Research Accomplishments.....	7
Reportable Outcomes.....	7
Conclusions.....	7
References.....	7
Appendices.....	

Introduction: Understanding the mechanisms involved in the progression from carcinoma *in situ* to metastatic disease is one of the most important problems in breast cancer research. In simple terms, progression to metastatic disease requires that breast carcinoma cells acquire the abilities to invade, stimulate angiogenesis and survive in non-breast tissues. Recently, progress has been made in identifying specific molecules that contribute to each of these critical events. These molecules include growth and angiogenic factors, as well as their corresponding receptors, and cell adhesion molecules. The challenge ahead is to understand how these molecules cooperate and interact to promote progression. Studies by our group and others have implicated a critical role for the $\alpha 6$ integrins in breast cancer progression (1) (2-4) (5). The goal of this IDEA proposal is to define the mechanism by which an integrin, such as the $\alpha 6$ integrins contributes to the survival of metastatic breast carcinoma cells. We postulated that these integrins regulate the expression of HIF-1 α , a key regulator of VEGF expression and angiogenesis in breast and other cancers. During the second year of this grant, we have made considerable progress that substantiates our hypothesis and completing our statement of work. We developed an effective RNAi strategy for inhibiting the expression of $\alpha 6$ integrins in breast carcinoma cells and used this approach to demonstrate that these integrins are necessary for breast carcinoma survival. Importantly, we have also obtained key data that the $\alpha 6$ integrins are essential for the transcriptional activation of HIF-1 α .

Body:

Development of an effective RNAi strategy for inhibiting expression of $\alpha 6$ integrins in breast carcinoma cells: The use of RNAi enables us to reduce expression of the $\alpha 6$ integrins specifically and to assess the consequences on breast carcinoma function. To this end, we designed oligonucleotides for this purpose and expressed them in MDA-435 cells. As shown in Fig. 1 below, this approach resulted in a significant decrease in the expression of the $\alpha 6\beta 1$ integrin but not of the $\alpha 5\beta 1$ integrin. The success of this approach allowed us to use it in subsequent experiments aimed at understanding the contribution of $\alpha 6\beta 1$ to VEGF transcription and breast carcinoma survival.

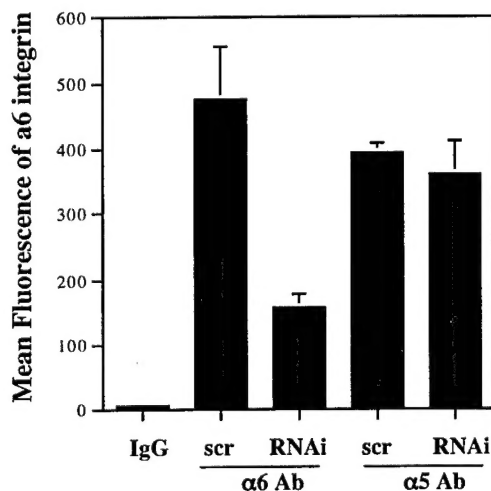
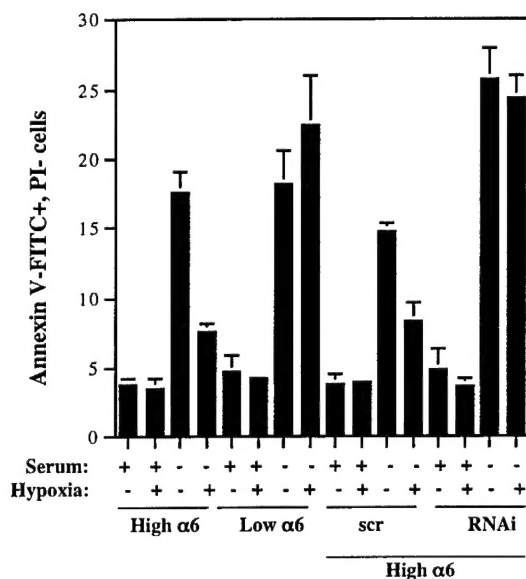


Figure 1: Use of RNAi to Reduce $\alpha 6\beta 1$ Integrin Expression in Breast Carcinoma Cells. Oligonucleotides were designed that inhibit expression of the $\alpha 6$ integrin subunit. These oligos, as well as a scrambled control sequence, were transfected into MDA-435 cells. After 72 hrs, the surface expression of the $\alpha 6$ and $\alpha 5$ integrin subunits was assessed by flow cytometry. Note that the expression of $\alpha 6$ but not $\alpha 5$ is reduced substantially.

Loss of $\alpha 6 \beta 1$ Integrin Expression Increases Apoptosis of Breast Carcinoma Cells in Stress Conditions: To examine the hypothesis that $\alpha 6 \beta 1$ integrin promotes breast carcinoma cell survival under stress condition, we incubated MDA-435 cells in either normoxia or hypoxia in low serum (0.5% FBS) for 24 hours and then measured annexin-FITC staining (Fig. 2). Serum deprivation dramatically increased apoptosis up to 4 fold while presence of 10% FBS protects cells from apoptosis regardless of $\alpha 6$ integrin level (Fig. 2). However, in high $\alpha 6$ clone, hypoxia reduced the apoptotic level approximately 50% suggesting that hypoxia acts as a survival factor (Fig. 2). Hypoxia driven cell survival against serum starvation is $\alpha 6 \beta 1$ integrin dependent because low $\alpha 6$ clone and $\alpha 6$ RNAi treated MDA-435 cells remained highly apoptotic under hypoxia (Fig. 2).



3. The $\alpha 6 \beta 1$ Integrin Regulates HIF-1 α Activation in Breast Carcinoma Cells: Since HIF-1 has been reported to play a major role in target gene transcription under hypoxia, we assessed the induction of HIF-1 α expression by hypoxia in clones of MDA-435 cells that exhibited either low or high $\alpha 6 \beta 1$ levels. As shown in Fig. 3, HIF-1 α induction was comparable on these clones, suggesting that the $\alpha 6 \beta 1$ integrin is not involved in the regulation of expression or stabilization of HIF-1 α .

Blot

HIF-1 α

	N	6hr	24hr	N	6hr	24hr
the	Hypoxia			Hypoxia		
a	high $\alpha 6$ clone			low $\alpha 6$ clone		

Figure 3. HIF-1 α Induction by Hypoxia is Independent of $\alpha 6 \beta 1$ Integrin Levels. MDA-435 breast carcinoma cells that exhibited either high or low expression of $\alpha 6 \beta 1$ were incubated in either normoxia or hypoxia for indicated times. Detergent extracts were blotted with HIF-1 α specific Ab.

To assess HIF-1 α activation, we monitored the association of HIF-1 with its co-activator, p300 (Fig. 4). HIF-1 binding to HRE itself is not enough to activate target gene transcription and HIF-1 association with p300 is required to recruit DNA pol II complex to initiate transcription. Therefore, the association of HIF-1 with p300 is an indicator of its activation. The association of HIF-1 with p300 was maximal level at 6 hour and gradually declined in high $\alpha 6$ clone (Fig. 5B). In the low $\alpha 6$ clone, the association of HIF-1 with p300 was barely detectable (Fig. 5B). The level of p300 expression remained constant in normoxia and hypoxia in both the high and low high $\alpha 6$ clones (Fig. 5A and B). The results indicate that, although the induction of HIF-1 α is comparable during hypoxia regardless of $\alpha 6\beta 1$ integrin level, $\alpha 6\beta 1$ integrin is involved in HIF-1 activation.

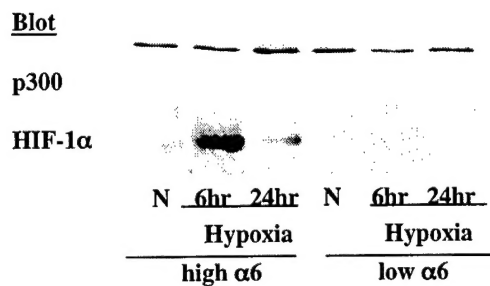


Figure 4. The $\alpha 6\beta 1$ Integrin is Necessary for HIF-1 α Activation in Breast Carcinoma Cells. MDA-435 breast carcinoma cells were prepared as described in Fig. 3, immunoprecipitated with a p300Ab and blotted with a HIF-1 α Ab.

Finally, to confirm the role of HIF-1 α in VEGF transcription under hypoxia, we used a HIF-1 α dominant negative mutant that blocks HIF-1 α binding to HRE. Expression of two different dominant negative HIF-1 α mutant blocked activation of VEGF transcription as shown by luciferase reading (Fig. 5), indicating that HIF-1 α and not other transcription factors, is playing a major role in VEGF transcription under hypoxia. Taken together with the data shown in Figs 2-4, these results provide direct evidence for the involvement of $\alpha 6\beta 1$ in HIF-1 α -mediated VEGF transcription.

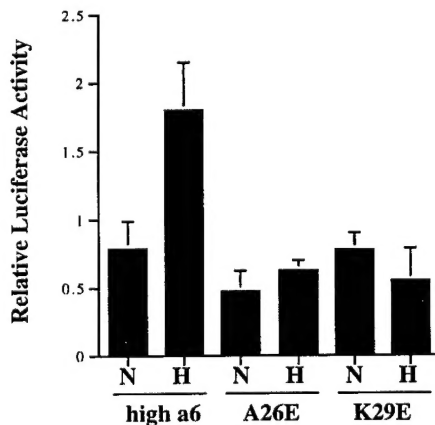


Figure 5. VEGF Transcription is HIF-1 α Dependent in Breast Carcinoma Cells. MDA-435 cells that expressed high levels of the $\alpha 6\beta 1$ integrin were transfected with either a control vector (high $\alpha 6$) or one of two dominant negative HIF-1 α constructs. Cells were maintained in either normoxia or hypoxia for 24 hrs and VEGF transcription was assessed by luciferase assay.

Key Research Accomplishments:

- Developed effective RNAi strategy for inhibiting expression of $\alpha 6$ integrins in breast carcinoma cells.
- Demonstrated the importance of $\alpha 6\beta 1$ integrin for the survival of breast carcinoma cells in hypoxia using this RNAi strategy.
- Established that $\alpha 6\beta 1$ regulates the activation of HIF-1 α in hypoxia but not its expression.
- Provided evidence that VEGF transcription in breast carcinoma cells is HIF-1 α dependent.

Reportable Outcomes:

A major manuscript that includes the results described in this report is currently in preparation and will be completed during the first part of Year 3 of this award.

Conclusions:

During Year 2 of this award, we have completed a significant amount of original Statement of Work. Most importantly, the data we have obtained validate the central hypothesis of our original proposal. In the remaining year of the award, we will complete the remaining items in Tasks 2 and 3 of our SOW.

References:

1. Friedrichs, K., Ruiz, P., Franke, F., Gille, I., Terpe, H. J., and Imhof, B. A. (1995) High expression level of alpha 6 integrin in human breast carcinoma is correlated with reduced survival. *Cancer Res* 55, 901-906
2. Shaw, L. M., Chao, C., Wewer, U. M., and Mercurio, A. M. (1996) Function of the integrin alpha 6 beta 1 in metastatic breast carcinoma cells assessed by expression of a dominant-negative receptor. *Cancer Res* 56, 959-963
3. Shaw, L. M., Rabinovitz, I., Wang, H. H., Toker, A., and Mercurio, A. M. (1997) Activation of phosphoinositide 3-OH kinase by the alpha6beta4 integrin promotes carcinoma invasion. *Cell* 91, 949-960
4. Wewer, U. M., Shaw, L. M., Albrechtsen, R., and Mercurio, A. M. (1997) The integrin alpha 6 beta 1 promotes the survival of metastatic human breast carcinoma cells in mice. *Am J Pathol* 151, 1191-1198
5. Mukhopadhyay, R., Theriault, R. L., and Price, J. E. (1999) Increased levels of alpha6 integrins are associated with the metastatic phenotype of human breast cancer cells. *Clin Exp Metastasis* 17, 325-332

Appendix:

None